

10/581,499

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number searching
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enhanced
NEWS 6 OCT 22 WPIDS, WPINDEX, and WPIX enhanced with Canadian PCT
Applications
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searching
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NEWS EXPRESS JUNE 27 08 CURRENT WINDOWS VERSION IS V8.3,
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FILE 'HOME' ENTERED AT 12:36:24 ON 06 DEC 2008

=> file reg

COST IN U.S. DOLLARS

SINCE FILE

TOTAL

10/581,499

	ENTRY	SESSION
FULL ESTIMATED COST	0.21	0.21

FILE 'REGISTRY' ENTERED AT 12:36:41 ON 06 DEC 2008
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STRUCTURE FILE UPDATES: 5 DEC 2008 HIGHEST RN 1080697-25-1
DICTIONARY FILE UPDATES: 5 DEC 2008 HIGHEST RN 1080697-25-1

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TSCA INFORMATION NOW CURRENT THROUGH July 5, 2008.

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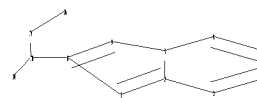
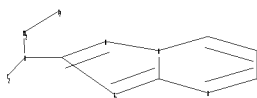
REGISTRY includes numerically searchable data for experimental and
predicted properties as well as tags indicating availability of
experimental property data in the original document. For information
on property searching in REGISTRY, refer to:

<http://www.cas.org/support/stngen/stndoc/properties.html>

=>

Uploading C:\Program Files\Stnexp\Queries\10581499.str

10/581,499



```
chain nodes :
10 14 15 16
ring nodes :
1 2 3 4 5 6 7 8 9
chain bonds :
8-10 10-14 10-15 15-16
ring bonds :
1-2 1-6 2-3 2-7 3-4 3-9 4-5 5-6 7-8 8-9
exact/norm bonds :
1-2 1-6 2-3 2-7 3-4 3-9 4-5 5-6 7-8 8-9 8-10 10-14 10-15 15-16
isolated ring systems :
containing 1 :
```

G1:C,N

G2:Ak,H

```
Match level :
1:Atom 2:Atom 3:Atom 4:Atom 5:Atom 6:Atom 7:Atom 8:Atom 9:Atom 10:CLASS
14:CLASS 15:CLASS 16:Atom
Generic attributes :
16:
Number of Carbon Atoms : less than 7
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Type of Ring System : Monocyclic

Element Count :
Node 16: Limited

C,C4-6

O,O0

S,S0

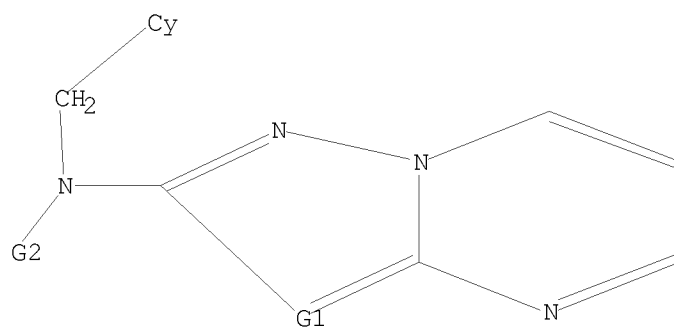
N,N0-2

L1 STRUCTURE UPLOADED

=> d l1

L1 HAS NO ANSWERS

L1 STR



G1 C,N

G2 Ak,H

Structure attributes must be viewed using STN Express query preparation.

=> s l1 sss sam

SAMPLE SEARCH INITIATED 12:37:08 FILE 'REGISTRY'

SAMPLE SCREEN SEARCH COMPLETED - 284 TO ITERATE

100.0% PROCESSED 284 ITERATIONS

6 ANSWERS

SEARCH TIME: 00.00.01

FULL FILE PROJECTIONS: ONLINE **COMPLETE**

BATCH **COMPLETE**

PROJECTED ITERATIONS: 4669 TO 6691

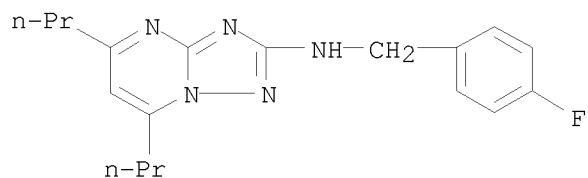
PROJECTED ANSWERS: 6 TO 266

L2 6 SEA SSS SAM L1

=> d scan l2

10/581,499

L2 6 ANSWERS REGISTRY COPYRIGHT 2008 ACS on STN
IN [1,2,4]Triazolo[1,5-a]pyrimidin-2-amine,
N-[(4-fluorophenyl)methyl]-5,7-dipropyl-
MF C18 H22 F N5

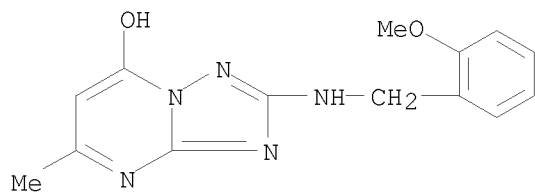


PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

HOW MANY MORE ANSWERS DO YOU WISH TO SCAN? (1):5

10/581,499

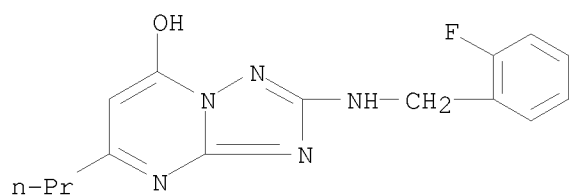
L2 6 ANSWERS REGISTRY COPYRIGHT 2008 ACS on STN
IN INDEX NAME NOT YET ASSIGNED
MF C14 H15 N5 O2



PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

10/581,499

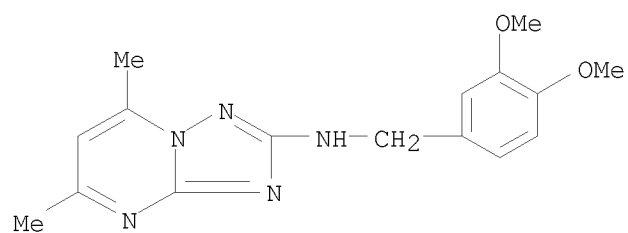
L2 6 ANSWERS REGISTRY COPYRIGHT 2008 ACS on STN
IN INDEX NAME NOT YET ASSIGNED
MF C15 H16 F N5 O



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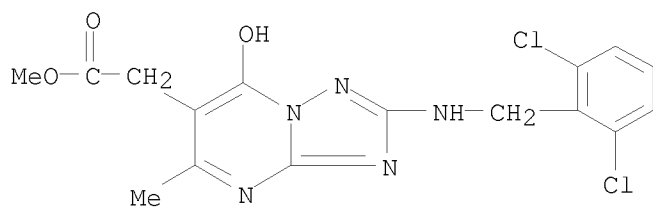
10/581,499

L2 6 ANSWERS REGISTRY COPYRIGHT 2008 ACS on STN
IN [1,2,4]Triazolo[1,5-a]pyrimidin-2-amine,
N-[(3,4-dimethoxyphenyl)methyl]-5,7-dimethyl-
MF C16 H19 N5 O2



10/581,499

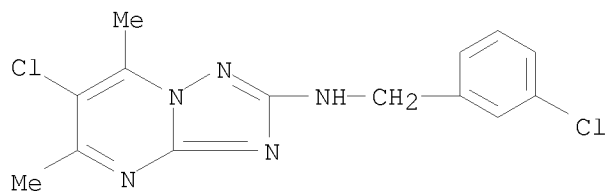
L2 6 ANSWERS REGISTRY COPYRIGHT 2008 ACS on STN
IN [1,2,4]Triazolo[1,5-a]pyrimidine-6-acetic acid,
2-[[(2,6-dichlorophenyl)methyl]amino]-7-hydroxy-5-methyl-, methyl ester
MF C16 H15 Cl2 N5 O3



PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

10/581,499

L2 6 ANSWERS REGISTRY COPYRIGHT 2008 ACS on STN
IN [1,2,4]Triazolo[1,5-a]pyrimidin-2-amine,
6-chloro-N-[(3-chlorophenyl)methyl]-5,7-dimethyl-
MF C14 H13 Cl2 N5



ALL ANSWERS HAVE BEEN SCANNED

10/581,499

=> s l1 sss ful
FULL SEARCH INITIATED 12:37:38 FILE 'REGISTRY'
FULL SCREEN SEARCH COMPLETED - 5508 TO ITERATE

100.0% PROCESSED 5508 ITERATIONS 158 ANSWERS
SEARCH TIME: 00.00.01

L3 158 SEA SSS FUL L1

=> file caplus
COST IN U.S. DOLLARS SINCE FILE TOTAL
ENTRY SESSION
FULL ESTIMATED COST 178.82 179.03

FILE 'CAPLUS' ENTERED AT 12:37:44 ON 06 DEC 2008
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FILE COVERS 1907 - 6 Dec 2008 VOL 149 ISS 24
FILE LAST UPDATED: 5 Dec 2008 (20081205/ED)

Caplus now includes complete International Patent Classification (IPC) reclassification data for the third quarter of 2008.

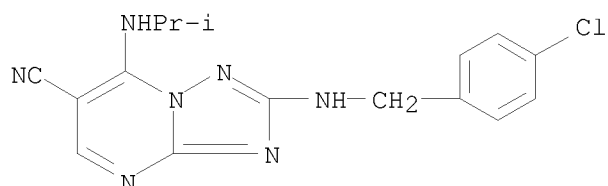
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<http://www.cas.org/legal/infopolicy.html>

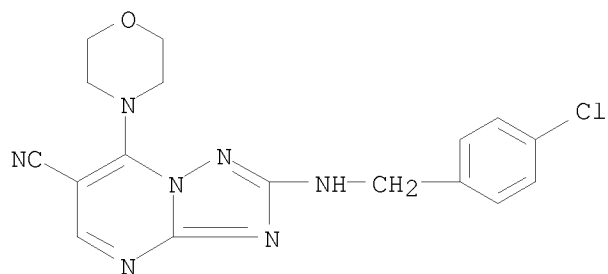
=> s l3
L4 10 L3

=> d l4 1-10 bib hitstr

L4 ANSWER 1 OF 10 CAPLUS COPYRIGHT 2008 ACS on STN
 AN 2007:1293045 CAPLUS
 DN 148:121662
 TI Preparation of 2,7-diamino-substituted
 [1,2,4]triazolo[1,5-a]pyrimidine-6-carbonitriles by solid-phase synthesis
 AU Cavallaro, Cullen L.; Harikrishnan, Lalgudi S.; Chi, Feng; Dodd, Dharmpal;
 Purandare, Ashok
 CS R & D, Bristol-Myers Squibb, Princeton, NJ, 08540, USA
 SO Journal of Combinatorial Chemistry (2008), 10(1), 28-30
 CODEN: JCCHFF; ISSN: 1520-4766
 PB American Chemical Society
 DT Journal
 LA English
 OS CASREACT 148:121662
 IT 1000982-70-6P 1000982-71-7P 1000982-72-8P
 RL: SPN (Synthetic preparation); PREP (Preparation)
 (solid-phase synthesis of diamino-substituted
 [1,2,4]triazolo[1,5-a]pyrimidinecarbonitriles)
 RN 1000982-70-6 CAPLUS
 CN [1,2,4]Triazolo[1,5-a]pyrimidine-6-carbonitrile,
 2-[[(4-chlorophenyl)methyl]amino]-7-[(1-methylethyl)amino]- (CA INDEX
 NAME)

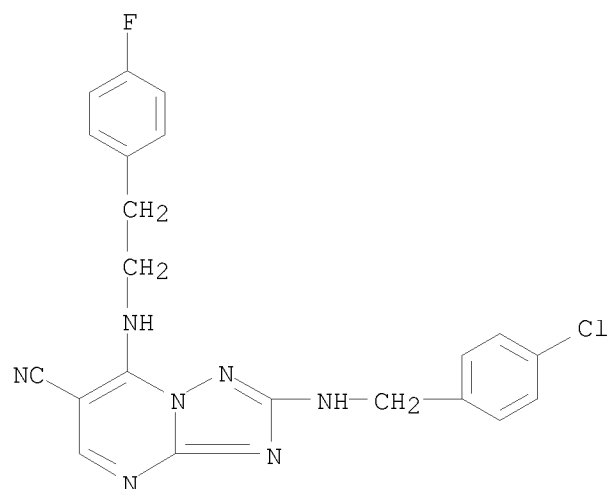


RN 1000982-71-7 CAPLUS
 CN [1,2,4]Triazolo[1,5-a]pyrimidine-6-carbonitrile,
 2-[[(4-chlorophenyl)methyl]amino]-7-(4-morpholinyl)- (CA INDEX NAME)



RN 1000982-72-8 CAPLUS
 CN [1,2,4]Triazolo[1,5-a]pyrimidine-6-carbonitrile,
 2-[[(4-chlorophenyl)methyl]amino]-7-[[2-(4-fluorophenyl)ethyl]amino]- (CA
 INDEX NAME)

10/581,499

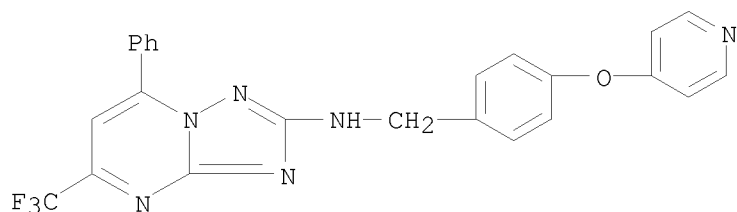


RE.CNT 37 THERE ARE 37 CITED REFERENCES AVAILABLE FOR THIS RECORD
ALL CITATIONS AVAILABLE IN THE RE FORMAT

L4 ANSWER 2 OF 10 CAPLUS COPYRIGHT 2008 ACS on STN
 AN 2005:523459 CAPLUS
 DN 143:60006
 TI Preparation of triazolo[1,5-a]pyrimidines and related compounds as TIE-2
 kinase inhibitors
 IN Schiemann, Kai; Hoelzemann, Guenter; Rautenberg, Wilfried
 PA Merck Patent G.m.b.H., Germany
 SO PCT Int. Appl., 188 pp.
 CODEN: PIXXD2
 DT Patent
 LA German
 FAN.CNT 1

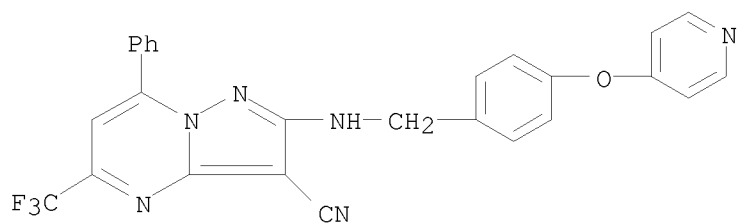
	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 2005054246	A2	20050616	WO 2004-EP12523	20041105
	WO 2005054246	A3	20050728		
	W:	AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NA, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW			
	RW:	BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LU, MC, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG			
	DE 10356579	A1	20050707	DE 2003-10356579	20031204
	AU 2004295032	A1	20050616	AU 2004-295032	20041105
	CA 2548156	A1	20050616	CA 2004-2548156	20041105
	EP 1727820	A2	20061206	EP 2004-797640	20041105
	R:	AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LI, LU, MC, NL, PL, PT, RO, SE, SI, SK, TR			
	JP 2007513099	T	20070524	JP 2006-541813	20041105
	US 20070112006	A1	20070517	US 2006-581499	20060602
PRAI	DE 2003-10356579	A	20031204		
	WO 2004-EP12523	W	20041105		
OS	MARPAT 143:60006				
IT	854272-88-1P 854272-95-0P 854273-14-6P 854273-39-5P 854273-42-0P 854273-43-1P 854273-56-6P 854273-62-4P 854273-70-4P 854273-73-7P 854274-25-2P 854274-26-3P 854274-58-1P				
	RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)				
	(preparation of triazolo[1,5-a]pyrimidines and related compds. as TIE-2 kinase inhibitors)				
RN	854272-88-1 CAPLUS				
CN	[1,2,4]Triazolo[1,5-a]pyrimidin-2-amine, 7-phenyl-N-[[4-(4-pyridinyloxy)phenyl]methyl]-5-(trifluoromethyl)- (CA INDEX NAME)				

10/581,499



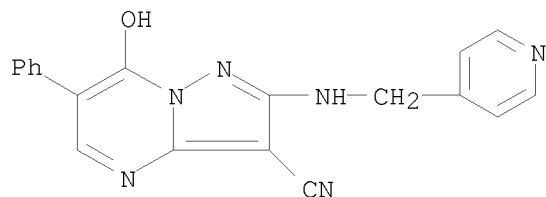
RN 854272-95-0 CAPLUS

CN Pyrazolo[1,5-a]pyrimidine-3-carbonitrile,
7-phenyl-2-[[4-(4-pyridinyloxy)phenyl]methyl]amino]-5-(trifluoromethyl)-
(CA INDEX NAME)



RN 854273-14-6 CAPLUS

CN Pyrazolo[1,5-a]pyrimidine-3-carbonitrile,
7-hydroxy-6-phenyl-2-[(4-pyridinylmethyl)amino]-, hydrochloride (1:1) (CA
INDEX NAME)

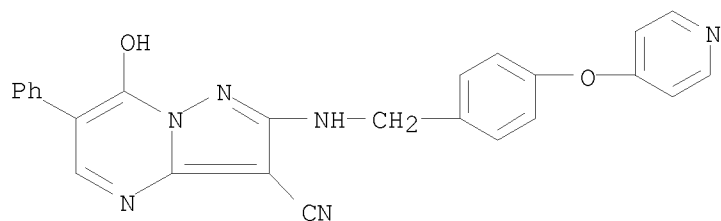


● HCl

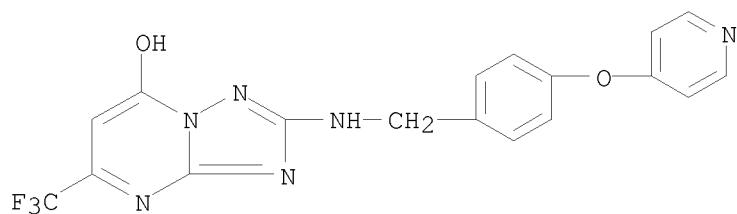
RN 854273-39-5 CAPLUS

CN Pyrazolo[1,5-a]pyrimidine-3-carbonitrile,
7-hydroxy-6-phenyl-2-[[[4-(4-pyridinyloxy)phenyl]methyl]amino]- (CA INDEX
NAME)

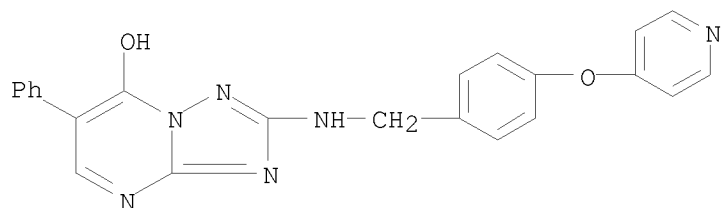
10/581,499



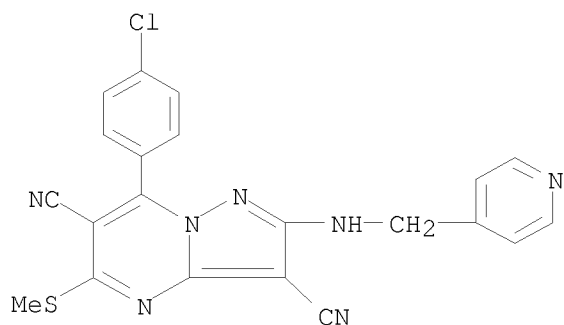
RN 854273-42-0 CAPLUS
CN [1,2,4]Triazolo[1,5-a]pyrimidin-7-ol,
2-[[[4-(4-pyridinyloxy)phenyl]methyl]amino]-5-(trifluoromethyl)- (CA
INDEX NAME)



RN 854273-43-1 CAPLUS
CN [1,2,4]Triazolo[1,5-a]pyrimidin-7-ol,
6-phenyl-2-[[[4-(4-pyridinyloxy)phenyl]methyl]amino]- (CA INDEX NAME)

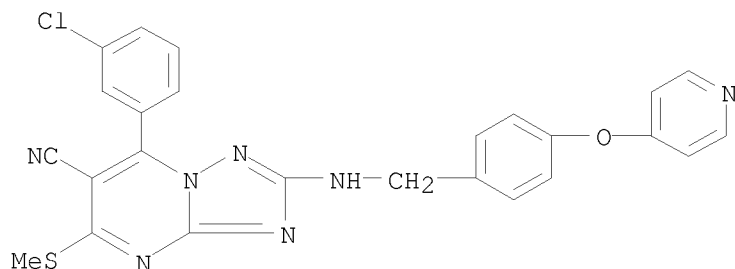


RN 854273-56-6 CAPLUS
CN Pyrazolo[1,5-a]pyrimidine-3,6-dicarbonitrile,
7-(4-chlorophenyl)-5-(methylthio)-2-[(4-pyridinylmethyl)amino]- (CA INDEX
NAME)

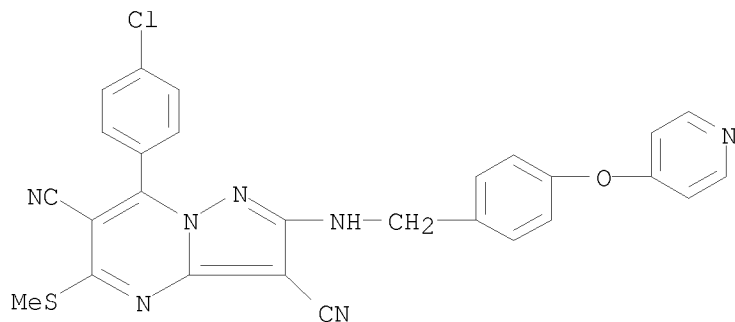


10/581,499

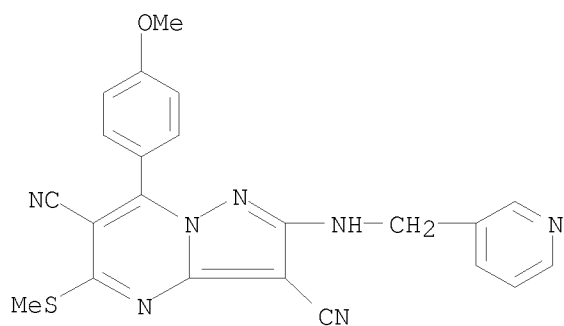
RN 854273-62-4 CAPLUS
CN [1,2,4]Triazolo[1,5-a]pyrimidine-6-carbonitrile,
7-(3-chlorophenyl)-5-(methylthio)-2-[[[4-(4-
pyridinyloxy)phenyl]methyl]amino]- (CA INDEX NAME)



RN 854273-70-4 CAPLUS
CN Pyrazolo[1,5-a]pyrimidine-3,6-dicarbonitrile,
7-(4-chlorophenyl)-5-(methylthio)-2-[[[4-(4-
pyridinyloxy)phenyl]methyl]amino]- (CA INDEX NAME)

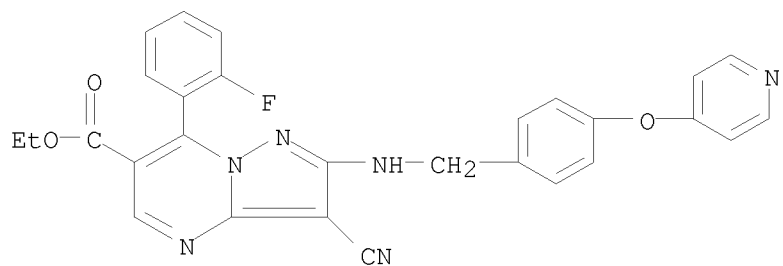


RN 854273-73-7 CAPLUS
CN Pyrazolo[1,5-a]pyrimidine-3,6-dicarbonitrile,
7-(4-methoxyphenyl)-5-(methylthio)-2-[(3-pyridinylmethyl)amino]- (CA
INDEX NAME)

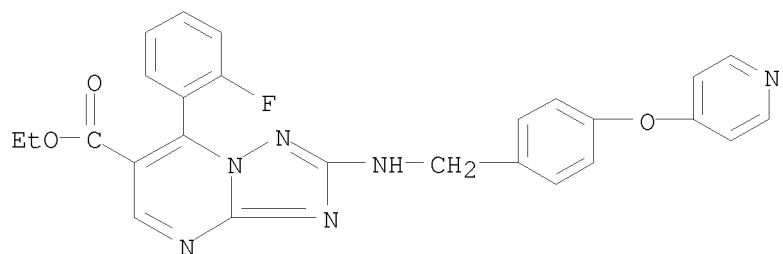


RN 854274-25-2 CAPLUS
CN Pyrazolo[1,5-a]pyrimidine-6-carboxylic acid,
3-cyano-7-(2-fluorophenyl)-2-[[[4-(4-pyridinyloxy)phenyl]methyl]amino]-,
ethyl ester (CA INDEX NAME)

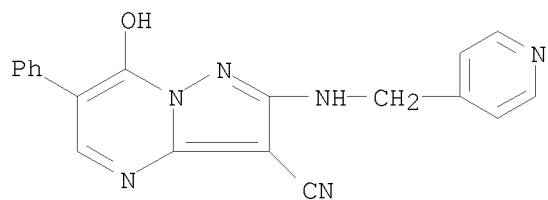
10/581,499



RN 854274-26-3 CAPLUS
CN [1,2,4]Triazolo[1,5-a]pyrimidine-6-carboxylic acid,
7-(2-fluorophenyl)-2-[[[4-(4-pyridinyloxy)phenyl]methyl]amino]-, ethyl
ester (CA INDEX NAME)



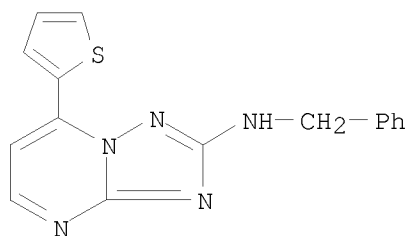
RN 854274-58-1 CAPLUS
CN Pyrazolo[1,5-a]pyrimidine-3-carbonitrile,
7-hydroxy-6-phenyl-2-[(4-pyridinylmethyl)amino]- (CA INDEX NAME)



10/581,499

L4 ANSWER 3 OF 10 CAPLUS COPYRIGHT 2008 ACS on STN
AN 2005:371263 CAPLUS
DN 142:430291
TI Preparation of substituted 2-amino-[1,2,4]triazolo[1,5-a]pyrimidine
derivatives as immunosuppressants
IN Kuramochi, Hiroshi; Masuda, Akira; Shimizu, Kazuhisa; Toyoda, Eriko;
Tokunaka, Kazuhiro
PA Nippon Kayaku Kabushiki Kaisha, Japan
SO PCT Int. Appl., 67 pp.
CODEN: PIXXD2
DT Patent
LA Japanese
FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 2005037837	A1	20050428	WO 2004-JP15245	20041015
	W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NA, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW				
	RW: BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG				
	CA 2542290	A1	20050428	CA 2004-2542290	20041015
	EP 1674454	A1	20060628	EP 2004-792462	20041015
	R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, FI, RO, CY, TR, BG, CZ, EE, HU, PL, SK				
	US 20070129383	A1	20070607	US 2006-575527	20060530
PRAI	JP 2003-357143	A	20031017		
	WO 2004-JP15245	W	20041015		
OS	MARPAT 142:430291				
IT	850733-95-8P				
	RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)				
	(preparation of substituted 2-amino-[1,2,4]triazolo[1,5-a]pyrimidine derivs. as immunosuppressants)				
RN	850733-95-8 CAPLUS				
CN	[1,2,4]Triazolo[1,5-a]pyrimidin-2-amine, N-(phenylmethyl)-7-(2-thienyl)- (CA INDEX NAME)				



RE.CNT 12 THERE ARE 12 CITED REFERENCES AVAILABLE FOR THIS RECORD
ALL CITATIONS AVAILABLE IN THE RE FORMAT

L4 ANSWER 4 OF 10 CAPLUS COPYRIGHT 2008 ACS on STN

AN 2005:33492 CAPLUS

DN 142:127563

TI Pyrazolo[1,5-a]pyrimidine derivatives, prophylactic or therapeutic agents containing them for protein tyrosine kinase-related diseases, and combination drugs containing them

IN Mukoyama, Harunobu; Shiohara, Hiroaki; Nishimura, Toshihiro; Nakayama, Akiko; Kikuchi, Shinji; Komatsu, Yoshimitsu; Onoda, Hideki

PA Kissei Pharmaceutical Co., Ltd., Japan

SO Jpn. Kokai Tokkyo Koho, 80 pp.

CODEN: JKXXAF

DT Patent

LA Japanese

FAN.CNT 1

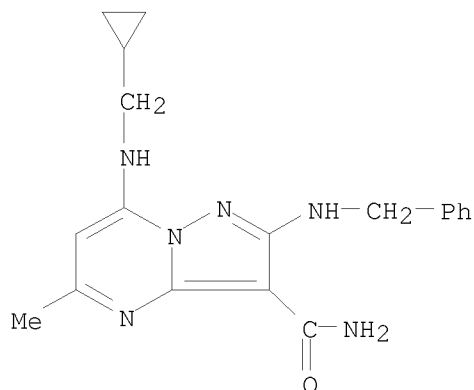
	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
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PI	JP 2005008581	A	20050113	JP 2003-175930	20030620
PRAI	JP 2003-175930		20030620		
OS	MARPAT 142:127563				
IT	824400-53-5P				

RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

(preparation of pyrazolopyrimidine derivs. as tyrosine kinase inhibitors for prevention and treatment of cancer, bone diseases, parkinsonism, GVHD, etc.)

RN 824400-53-5 CAPLUS

CN Pyrazolo[1,5-a]pyrimidine-3-carboxamide,
7-[(cyclopropylmethyl)amino]-5-methyl-2-[(phenylmethyl)amino]- (CA INDEX NAME)



IT 824398-07-4P 824399-31-7P

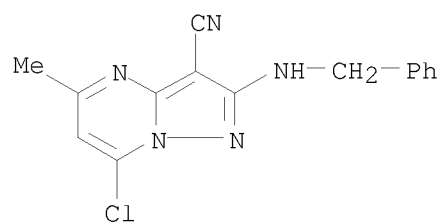
RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)

(preparation of pyrazolopyrimidine derivs. as tyrosine kinase inhibitors for prevention and treatment of cancer, bone diseases, parkinsonism, GVHD, etc.)

RN 824398-07-4 CAPLUS

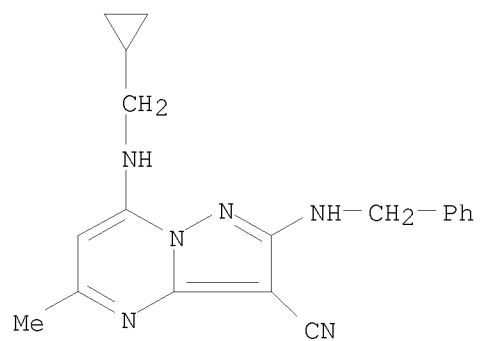
CN Pyrazolo[1,5-a]pyrimidine-3-carbonitrile,
7-chloro-5-methyl-2-[(phenylmethyl)amino]- (CA INDEX NAME)

10/581,499



RN 824399-31-7 CAPLUS

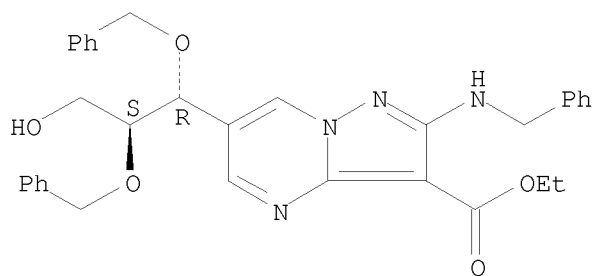
CN Pyrazolo[1,5-a]pyrimidine-3-carbonitrile,
7-[(cyclopropylmethyl)amino]-5-methyl-2-[(phenylmethyl)amino]- (CA INDEX
NAME)



10/581,499

L4 ANSWER 5 OF 10 CAPLUS COPYRIGHT 2008 ACS on STN
AN 2004:1102652 CAPLUS
DN 142:198293
TI Synthesis of heterocyclic compounds by ring transformations of 2-formyl
pentose glycals
AU Bari, Ahmed; Feist, Holger; Michalik, Dirk; Michalik, Manfred; Peseke,
Klaus
CS Universitaet Rostock, Fachbereich Chemie, Rostock, 18051, Germany
SO Synthesis (2004), (17), 2863-2868
CODEN: SYNTBF; ISSN: 0039-7881
PB Georg Thieme Verlag
DT Journal
LA English
OS CASREACT 142:198293
IT 839716-48-2P
RL: SPN (Synthetic preparation); PREP (Preparation)
(synthesis of heterocyclic acyclo-C-nucleosides and open-chain
monosaccharide nucleoside analogs based on ring transformations of
2-formyl pentose glycals)
RN 839716-48-2 CAPLUS
CN Pyrazolo[1,5-a]pyrimidine-3-carboxylic acid,
6-[(1R,2S)-3-hydroxy-1,2-bis(phenylmethoxy)propyl]-2-[(phenylmethyl)amino]-
, ethyl ester (CA INDEX NAME)

Absolute stereochemistry. Rotation (-).



RE.CNT 19 THERE ARE 19 CITED REFERENCES AVAILABLE FOR THIS RECORD
ALL CITATIONS AVAILABLE IN THE RE FORMAT

L4 ANSWER 6 OF 10 CAPLUS COPYRIGHT 2008 ACS on STN
 AN 2002:637565 CAPLUS
 DN 137:185499
 TI Preparation of triazolopyrimidines as thrombin inhibitors
 IN Williams, Peter D.; Coburn, Craig; Burgey, Christopher; Morrisette, Matthew M.
 PA Merck & Co., Inc., USA
 SO PCT Int. Appl., 184 pp.
 CODEN: PIXXD2
 DT Patent
 LA English
 FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 2002064211	A1	20020822	WO 2002-US4654	20020205
	W:				
	AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, OM, PH, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZM, ZW				
	RW:				
	GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG				
	AU 2002247158	A1	20020828	AU 2002-247158	20020205
PRAI	US 2001-267813P	P	20010209		
	WO 2002-US4654	W	20020205		

OS MARPAT 137:185499

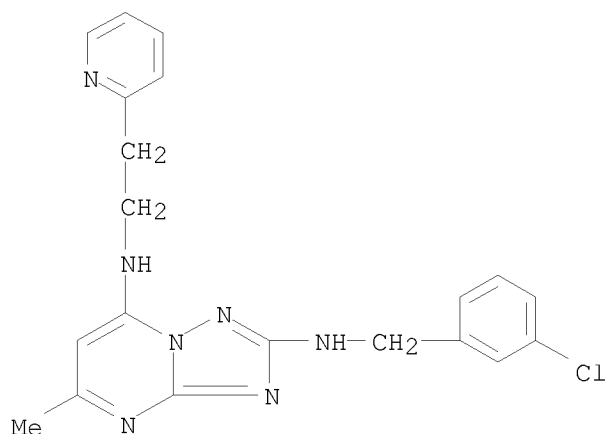
IT 450398-63-7P 450398-65-9P 450398-66-0P

RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

(claimed compound; preparation of triazolopyrimidines as thrombin inhibitors)

RN 450398-63-7 CAPLUS

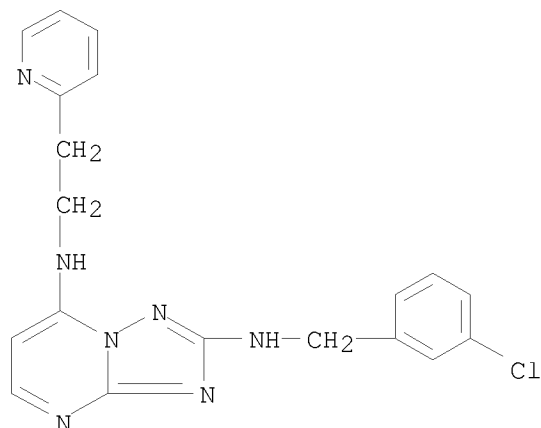
CN [1,2,4]Triazolo[1,5-a]pyrimidine-2,7-diamine,
 N2-[(3-chlorophenyl)methyl]-N7-[2-(2-pyridinyl)ethyl]- (CA INDEX NAME)



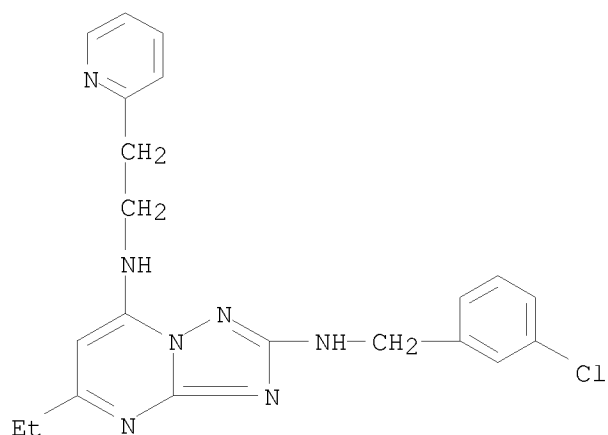
RN 450398-65-9 CAPLUS

CN [1,2,4]Triazolo[1,5-a]pyrimidine-2,7-diamine,
 N2-[(3-chlorophenyl)methyl]-N7-[2-(2-pyridinyl)ethyl]- (CA INDEX NAME)

10/581,499



RN 450398-66-0 CAPLUS
CN [1,2,4]Triazolo[1,5-a]pyrimidine-2,7-diamine,
N2-[(3-chlorophenyl)methyl]-5-ethyl-N7-[2-(2-pyridinyl)ethyl]- (CA INDEX
NAME)



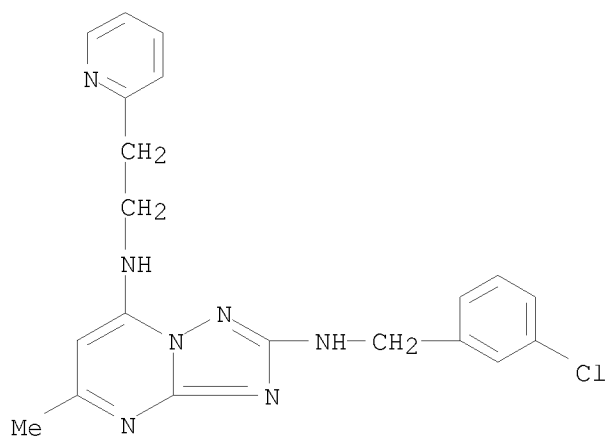
IT 450399-61-8P
RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU
(Therapeutic use); BIOL (Biological study); PREP (Preparation); USES
(Uses)
(preparation of triazolopyrimidines as thrombin inhibitors)

RN 450399-61-8 CAPLUS
CN [1,2,4]Triazolo[1,5-a]pyrimidine-2,7-diamine,
N2-[(3-chlorophenyl)methyl]-5-methyl-N7-[2-(2-pyridinyl)ethyl]-,
2,2,2-trifluoroacetate (20:29) (CA INDEX NAME)

CM 1

CRN 450398-63-7
CMF C20 H20 Cl N7

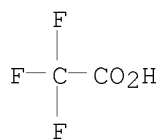
10/581,499



CM 2

CRN 76-05-1

CMF C2 H F3 O2



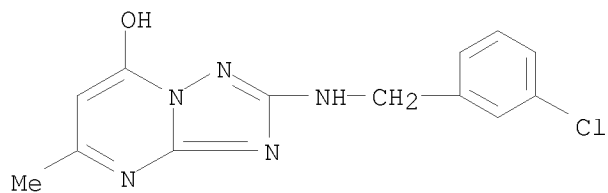
IT 450400-04-1P 450400-05-2P

RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)

(preparation of triazolopyrimidines as thrombin inhibitors)

RN 450400-04-1 CAPLUS

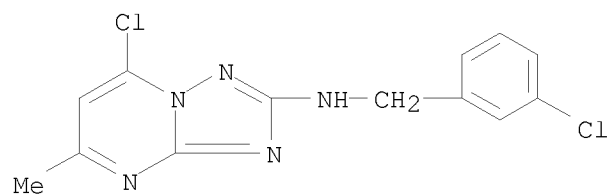
CN [1,2,4]Triazolo[1,5-a]pyrimidin-7-ol,
2-[[(3-chlorophenyl)methyl]amino]-5-methyl- (CA INDEX NAME)



RN 450400-05-2 CAPLUS

CN [1,2,4]Triazolo[1,5-a]pyrimidin-2-amine,
7-chloro-N-[(3-chlorophenyl)methyl]-5-methyl- (CA INDEX NAME)

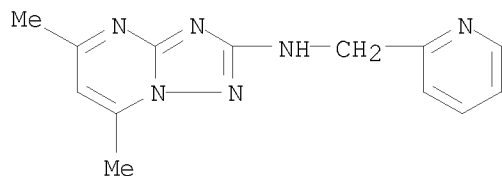
10/581,499



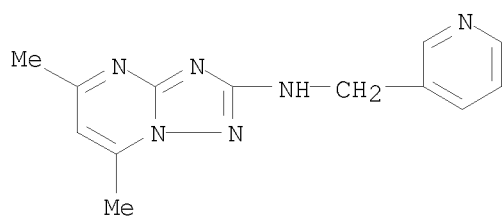
RE.CNT 4 THERE ARE 4 CITED REFERENCES AVAILABLE FOR THIS RECORD
ALL CITATIONS AVAILABLE IN THE RE FORMAT

10/581,499

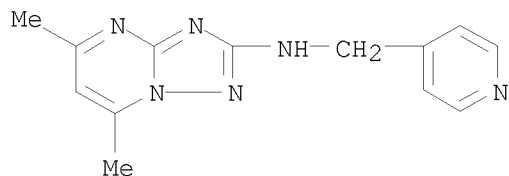
L4 ANSWER 7 OF 10 CAPLUS COPYRIGHT 2008 ACS on STN
AN 1995:548834 CAPLUS
DN 123:112014
OREF 123:20013a,20016a
TI On Triazoles. XXXV 1. The reaction of 5-amino-1,2,4-triazoles with di- and triketones
AU Reiter, Jozsef; Pongo, Laszlo; Koevesdi, Istvan; Pallagi, Istvan
CS EGIS Pharmaceuticals, Budapest, Hung.
SO Journal of Heterocyclic Chemistry (1995), 32(2), 407-17
CODEN: JHTCAD; ISSN: 0022-152X
PB HeteroCorporation
DT Journal
LA English
OS CASREACT 123:112014
IT 165684-56-0P 165684-57-1P 165684-58-2P
165684-75-3P 165684-76-4P 165684-77-5P
RL: SPN (Synthetic preparation); PREP (Preparation)
(preparation of)
RN 165684-56-0 CAPLUS
CN [1,2,4]Triazolo[1,5-a]pyrimidin-2-amine,
5,7-dimethyl-N-(2-pyridinylmethyl)- (CA INDEX NAME)



RN 165684-57-1 CAPLUS
CN [1,2,4]Triazolo[1,5-a]pyrimidin-2-amine,
5,7-dimethyl-N-(3-pyridinylmethyl)- (CA INDEX NAME)



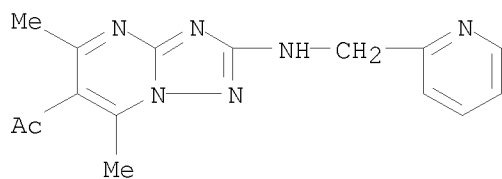
RN 165684-58-2 CAPLUS
CN [1,2,4]Triazolo[1,5-a]pyrimidin-2-amine,
5,7-dimethyl-N-(4-pyridinylmethyl)- (CA INDEX NAME)



RN 165684-75-3 CAPLUS

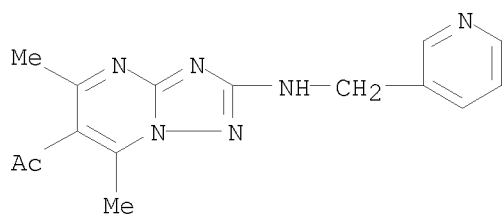
10/581,499

CN Ethanone, 1-[5,7-dimethyl-2-[(2-pyridinylmethyl)amino][1,2,4]triazolo[1,5-a]pyrimidin-6-yl]- (CA INDEX NAME)



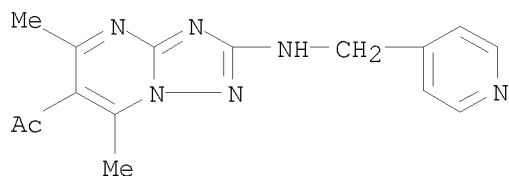
RN 165684-76-4 CAPLUS

CN Ethanone, 1-[5,7-dimethyl-2-[(3-pyridinylmethyl)amino][1,2,4]triazolo[1,5-a]pyrimidin-6-yl]- (CA INDEX NAME)



RN 165684-77-5 CAPLUS

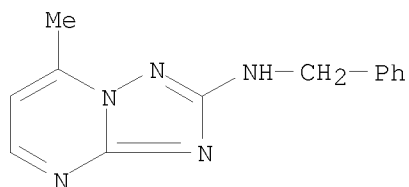
CN Ethanone, 1-[5,7-dimethyl-2-[(4-pyridinylmethyl)amino][1,2,4]triazolo[1,5-a]pyrimidin-6-yl]- (CA INDEX NAME)



10/581,499

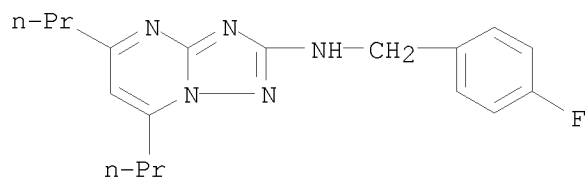
L4 ANSWER 8 OF 10 CAPLUS COPYRIGHT 2008 ACS on STN
AN 1991:656223 CAPLUS
DN 115:256223
OREF 115:43577a,43580a
TI Preparation of substituted 2-amino-1,2,4-triazolo[1,5-a]pyrimidines and
-triazines as herbicides
IN Wegner, Peter; Egner, Ursula; Saenger, Wolfram; Gerbling, Klaus Peter;
Johann, Gerhard; Rees, Richard
PA Schering A.-G., Germany
SO Ger. Offen., 13 pp.
CODEN: GWXXBX
DT Patent
LA German
FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
	-----	---	-----	-----	-----
PI	DE 4008181	A1	19910919	DE 1990-4008181	19900312
PRAI	DE 1990-4008181		19900312		
OS	CASREACT 115:256223; MARPAT 115:256223				
IT	137353-64-1P				
	RL: AGR (Agricultural use); BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); SPN (Synthetic preparation); BIOL (Biological study); PREP (Preparation); USES (Uses) (preparation of, as herbicide)				
RN	137353-64-1 CAPLUS				
CN	[1,2,4]Triazolo[1,5-a]pyrimidin-2-amine, 7-methyl-N-(phenylmethyl)- (CA INDEX NAME)				



10/581,499

L4 ANSWER 9 OF 10 CAPLUS COPYRIGHT 2008 ACS on STN
AN 1983:539891 CAPLUS
DN 99:139891
OREF 99:21493a,21496a
TI Dialkyl bicyclic heterocycles with a bridgehead nitrogen as purine analogs
possessing significant cardiac inotropic activity
AU Okabe, Takayuki; Bhooshan, Bharat; Novinson, Thomas; Hillyard, Ira W.;
Garner, Garland E.; Robins, Roland K.
CS Viratek, Inc., Covina, CA, 91732, USA
SO Journal of Heterocyclic Chemistry (1983), 20(3), 735-51
CODEN: JHTCAD; ISSN: 0022-152X
DT Journal
LA English
OS CASREACT 99:139891
IT 87253-54-1P
RL: BAC (Biological activity or effector, except adverse); BSU (Biological
study, unclassified); SPN (Synthetic preparation); BIOL (Biological
study); PREP (Preparation)
(preparation and inotropic activity of)
RN 87253-54-1 CAPLUS
CN [1,2,4]Triazolo[1,5-a]pyrimidin-2-amine,
N-[(4-fluorophenyl)methyl]-5,7-dipropyl- (CA INDEX NAME)



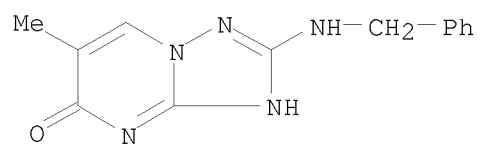
10/581,499

L4 ANSWER 10 OF 10 CAPLUS COPYRIGHT 2008 ACS on STN
AN 1970:425482 CAPLUS
DN 73:25482
OREF 73:4243a,4246a
TI Triazolo[1,5-a]pyrimidines
IN Dukes, Michael
PA Imperial Chemical Industries Ltd.
SO Ger. Offen., 75 pp.
CODEN: GWXXBX
DT Patent
LA German
FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
	-----	----	-----	-----	-----
PI	DE 1946315	A	19700319	DE 1969-1946315	19690912
	DE 1946315	C2	19850515		
	GB 1234635	A	19710609	GB 1968-43627	19680913
	ZA 6905832	A	19710331	ZA 1969-5832	19690814
	US 3689488	A	19720905	US 1969-850221	19690814
	PL 80261	B1	19750830	PL 1969-135379	19690815
	PL 80539	B1	19750830	PL 1969-139471	19690815
	PL 80662	B1	19750830	PL 1969-139525	19690815
	PL 80664	B1	19750830	PL 1969-139527	19690815
	DK 137498	C	19780828	DK 1969-4586	19690827
	BR 6912007	D0	19730419	BR 1969-212007	19690829
	SU 404249	A3	19731026	SU 1969-1447192	19690902
	SU 432719	A3	19740615	SU 1969-1445877	19690902
	SU 485597	A3	19750925	SU 1969-1447118	19690902
	SU 511001	A3	19760415	SU 1969-1365596	19690902
	CS 163197	B2	19750829	CS 1969-2756	19690903
	CS 163196	B2	19750829	CS 1969-6023	19690903
	BE 738830	A	19700312	BE 1969-738830	19690912
	NL 6913907	A	19700317	NL 1969-13907	19690912
	NL 162651	B	19800115		
	NL 162651	C	19800616		
	FR 2018077	A5	19700529	FR 1969-31200	19690912
	FR 2018077	B1	19730112		
	AT 292000	B	19710810	AT 1969-8717	19690912
	AT 292696	B	19710910	AT 1970-8667	19690912
	AT 292697	B	19710910	AT 1970-8668	19690912
	AT 292699	B	19710910	AT 1970-8670	19690912
	SE 373584	B	19750210	SE 1969-12601	19690912
	SE 377460	B	19750707	SE 1972-16479	19690912
	JP 51007677	B	19760310	JP 1969-72676	19690912
	ES 371509	A1	19711101	ES 1969-371509	19690913
	CH 522666	A	19720515	CH 1969-522666	19690915
	CH 523270	A	19720531	CH 1969-523270	19690915
	CH 523272	A	19720531	CH 1969-523272	19690915
	CH 529772	A	19721031	CH 1969-529772	19690915
	CH 530410	A	19721115	CH 1969-530410	19690915
	US 3773949	A	19731120	US 1972-252727	19720512
PRAI	GB 1968-43627	A	19680913		
	GB 1969-22266	A	19690501		
	US 1969-850221	A3	19690814		
	SU 1969-1365596	A	19690902		
IT	27276-78-4P				
	RL: SPN (Synthetic preparation); PREP (Preparation)				
	(preparation of)				
RN	27276-78-4 CAPLUS				
CN	[1,2,4]Triazolo[1,5-a]pyrimidin-5(1H)-one,				

10/581,499

6-methyl-2-[(phenylmethyl)amino]- (CA INDEX NAME)



10/581,499

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COST IN U.S. DOLLARS

SINCE FILE

TOTAL

ENTRY

SESSION

FULL ESTIMATED COST

37.98

217.01

STN INTERNATIONAL LOGOFF AT 12:38:12 ON 06 DEC 2008